

IN THE CLAIMS

Please amend the claims as follows.

Claims 1-52. Canceled.

- ~~531~~. (currently amended) A method of altering the binding affinity of a peptide to its receptor, comprising conjugating the peptide to an amphiphilic oligomer comprising a lipophilic moiety coupled to a hydrophilic moiety.
- ~~542~~. (currently amended) The method according to claim ~~481~~ further characterized in that the binding affinity is increased.
- ~~553~~. (currently amended) The method according to claim ~~481~~ further characterized in that the binding affinity is reduced.
- ~~564~~. (currently amended) The method of claim ~~481~~, wherein the peptide is a peptide or protein.
- ~~575~~. (currently amended) The method of claim ~~484~~, wherein the peptide is selected from the group consisting of: enkephalin, adrenocorticotrophic hormone, adenosine deaminase, ribonuclease, alkaline phosphatase, angiotensin, antibodies, arginase, arginine deaminase, asparaginase, caerulein, calcitonin, chemotrypsin, cholecystokinin, clotting factors, dynorphins, ~~endorphins~~, endorphins, ~~enkephalins~~, enkephalins, erythropoietin, gastrin-releasing peptide, glucagon, hemoglobin, hypothalamic releasing factors, interferon, katalcalcin, motilin, neuropeptide Y, ~~neuretensin~~neurotensin, non-naturally occurring opioids, ~~oxytocin~~oxytocin, papain, parathyroid hormone, ~~peptides~~-prolactin,

soluble CD-4, somatomedin, somatostatin, ~~somatostatin~~, somatotropin, superoxide dismutase, thyroid stimulating hormone, tissue plasminogen activator, trypsin, vasopressin, and analogues and fragments of such peptides.

~~586~~. (currently amended) The method of claim ~~484~~ wherein the peptide is [met⁵]enkephalin.

~~597~~. (currently amended) The method of claim ~~481~~, wherein the lipophilic moiety is selected from the group consisting of fatty acids, C₁₋₂₆alkyls, and ~~cholesterol~~cholesterol.

~~608~~. (currently amended) The method of claim ~~481~~, wherein the hydrophilic moiety is selected from the group consisting of sugars or PEG₁₋₇.

61-63. Canceled.

~~649~~. (new and currently amended) The method of claim 1, wherein the receptor is an opioid receptor.